Job Advertisement

Job Description: Senior Quality Manager - EU
Type: Full Time
Location: Belgium - Liege
Date: Immediate

About Telix Pharmaceuticals Limited

Telix Pharmaceuticals Limited ("Telix", the "Company") is an Australian public Company (ASX:TLX) headquartered in Melbourne with operations in Europe, the US and Japan. Our mission is to be a leading, global biopharmaceutical Company in the field of "theranostic" radiopharmaceuticals and the Company is currently developing a mid-late stage pipeline of products in oncology and rare diseases.

Description

Reporting to the Head of Quality – Europe, the Quality Manager will lead the quality management of the commercial scale-up following GMP requirements. As Senior QA Manager, you will hold Quality Assurance responsibility related to the Marketing Authorization holder and operations for one of Telix's key products. The presence onsite in Liège is largely required in the regulatory context and highly desirable from a management and collaboration standpoint. You will also have regular interaction with key staff in Australia and the USA.

Every day is different however core responsibilities will include:

- Management of the QMS and its continuous improvement, ensuring its application and follow up. Training, SOPs, reporting and internal quality reviews are all part of this remit.
- Reviewing documents, assessing risk and creating GAP analysis based on current regulations.
- Investigating and resolving product and process problems related to quality issues.
- Investigations of customer complaints and working with internal & external stakeholders to resolution.
- Ensuring CAPA's are properly documented and corrective actions taken
- Understand and adhere to GMP policies and procedures.
- Providing ongoing technical assistance and problem solving with existing manufacturers and suppliers, leveraging industry experience and best practices.

To be considered, you will require:

- A Bachelor's degree in relevant discipline.
- Demonstrated experience in a pharmaceutical manufacturing environment with Good Manufacturing Practices (GMP).
- Experience working in cross-functional teams in particular, R&D, Operations and Quality.
- Audit experience.
- Experience writing and reviewing SOPs and specifications.
- People management experience (direct reports) highly preferred.
- Languages: English and French fluency. Note: applications should be in English.

Agency support is not required for this role and no submissions will be considered.

For further details and information pertaining to compensation for the role, as well as expressions of interest, please contact Telix People and Culture at: employment@telixpharma.com